

STIC-Biotech/ChemLib

75096

Fr m: Portner, Ginny
Sent: Saturday, September 07, 2002 3:13 PM
T : STIC-Biotech/ChemLib
Subject: 09/674,254

Please search SEQ ID No 3 (peptide of 16 amino acids) .. thanks!

Ginny Portner
CM1, Art Unit 1645
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(703) 308-7543

RECEIVED
SEP-9 2002
(S113)

Point of Contact
Searcher: P. Sheppard
Phone: Telephone number: (703) 308-4499
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Date Picked Up: _____
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Online time: _____

TYPE OF SEARCH:
NA Sequences: _____
AA Sequences: _____
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST (where applic.)
STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: _____
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Other (specify): _____

WEST Search History

DATE: Saturday, September 07, 2002

<u>Set Name</u>	<u>Query</u>	<u>Hit Count</u>	<u>Set Name</u>
side by side			result set
<i>DB=USPT; PLUR=YES; OP=AND</i>			
L1	5916751.9n.	0	L1
L2	5916751.pn.	1	L2
L3	lefty.ti,ab,clm.	2	L3

END OF SEARCH HISTORY

WEST[Generate Collection](#)[Print](#)**Search Results - Record(s) 1 through 3 of 3 returned.**☐ 1. Document ID: US 6294662 B1

L3: Entry 1 of 3

File: USPT

Sep 25, 2001

DOCUMENT-IDENTIFIER: US 6294662 B1

TITLE: Nucleic acids encoding an endometrial bleeding associated factor (ebaf)

Abstract Text (1):

A method for the early diagnosing of selected adenocarcinomas in a human comprising the steps of removing a bodily sample from the human, and assaying the bodily sample for elevated expression of a specific gene. The gene being assayed for in the bodily sample is the TGFB-4 gene (hereinafter referred to as the endometrial bleeding associated factor (ebaf) gene). The bodily sample can be tissue from a specific organ in the body, or a blood sample. Increased levels of ebaf in the sample relative to basal levels may be indicative of a mucinous adenocarcinoma of the colon or ovaries, or an adenocarcinoma of the testis.

Brief Summary Text (22):

The specific gene referred to above is the TGFB-4 gene (hereinafter referred to as the endometrial bleeding associated factor (ebaf) gene). Applicants recently discovered this gene in humans (please see Ravi Kothapalli, Ibrahim Buyuksal, Shi-Qi Wu, Nasser Chegini, Siamak Tabibzadeh: Detection of ebaf, a novel human gene of the TGF- β superfamily; association of gene expression with endometrial bleeding J. Clin. Invest. 1997, 99:2342-2350, which is hereby incorporated by reference herein). The cDNA sequence of the ebaf gene is set forth in SEQ. ID NO. 1.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw Desc
Image												

☐ 2. Document ID: US 5916751 A

L3: Entry 2 of 3

File: USPT

Jun 29, 1999

DOCUMENT-IDENTIFIER: US 5916751 A

TITLE: Method for the diagnosis of selected adenocarcinomas**Abstract Text (1):**

A method for the early diagnosing of selected adenocarcinomas in a human comprising the steps of removing a bodily sample from the human, and assaying the bodily sample for elevated expression of a specific gene, the gene being assayed for in the bodily sample is the TGFB-4 gene (hereinafter referred to as the endometrial bleeding associated factor (ebaf) gene. The bodily sample can be tissue from a specific organ in the body, or a blood sample. Increased levels of ebaf in the sample relative to basal levels may be indicative of a mucinous adenocarcinoma of the colon or ovaries, or an adenocarcinoma of the testis.

Brief Summary Text (22):

The specific gene referred to above is the TGFB-4 gene (hereinafter referred to as the endometrial bleeding associated factor (ebaf) gene). Applicants recently discovered this gene in humans (please see, Ravi Kothapalli, Ibrahim Buyuksal, Shi-Qi Wu, Nasser Chegini, Siamak Tabibzadeh: Detection of ebaf, a novel human gene of the TGF-superfamily; association of gene expression with endometrial bleeding J. Clin. Invest. 1997, 99:2342-2350, which is hereby incorporated by reference herein). The cDNA sequence of the ebafgene is set forth in SEQ. ID NO. 1.

Other Reference Publication (6):

Siamak Tabibzadeh, Ravi Kothapalli, Ibrahim Buyuksal, "Distinct Tumor Specific Expression of TGFB4 (ebaf), a Novel Human Gene of the TGF-B Superfamily," Frontiers in Bioscience 2, Jul. 1997, pp. 18-25.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KIMC	Draw Desc
Image												

☐ **3. Document ID: US 5916751 A**

L3: Entry 3 of 3

File: DWPI

Jun 29, 1999

DERWENT-ACC-NO: 1999-384717

DERWENT-WEEK: 200175

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TITLE: Detecting serous or mucinous colon/ovarian adenocarcinomas and testicular adenocarcinoma by assaying for elevated expression of a gene

INVENTOR: KOTHAPALLI, R; TABIBZADEH, S

PRIORITY-DATA: 1996US-025800P (August 27, 1996), 1997US-0919421 (August 27, 1997)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
US 5916751 A	June 29, 1999		011	G01N033/574

INT-CL (IPC): C12 Q 1/68; G01 N 33/48; G01 N 33/574

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw Desc
Image												

[Generate Collection](#)[Print](#)

Terms	Documents
tgfb4 or tgfb-4	3

Display Format: [Change Format](#)[Previous Page](#)[Next Page](#)

WEST Search History

DATE: Saturday, September 07, 2002

<u>Set Name</u> side by side	<u>Query</u>	<u>Hit</u> <u>Count</u>	<u>Set</u> <u>Name</u> result set
<i>DB=USPT; PLUR=YES; OP=AND</i>			
L1	tgfb4 or tgfb-4	2	L1
<i>DB=USPT,PGPB,JPAB,EPAB,DWPI,TDBD; PLUR=YES; OP=AND</i>			
L2	L1	2	L2
L3	tgfb4 or tgfb-4	3	L3
L4	tgf-b4 or tgf-b-4	1	L4
L5	L4	1	L5

END OF SEARCH HISTORY

ile 155:MEDLINE(R) 1966-2002/Sep W1

*File 155: Alert feature enhanced for multiple files, duplicates removal, customized scheduling. See HELP ALERT.

Set Items Description

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?e transforming growth factor

Ref	Items	RT	Index-term
E1	4		TRANSFORMING ACIDIC COILED COIL PROTEIN
E2	0	1	TRANSFORMING GENES
E3	0		*TRANSFORMING GROWTH FACTOR
E4	2596	6	TRANSFORMING GROWTH FACTOR ALPHA
E5	20		TRANSFORMING GROWTH FACTOR ALPHA --ADMINISTRAT
E6	2		TRANSFORMING GROWTH FACTOR ALPHA --ADVERSE EFF
E7	1		TRANSFORMING GROWTH FACTOR ALPHA --AGONISTS --
E8	373		TRANSFORMING GROWTH FACTOR ALPHA --ANALYSIS --
E9	29		TRANSFORMING GROWTH FACTOR ALPHA --ANTAGONISTS
E10	331		TRANSFORMING GROWTH FACTOR ALPHA --BIOSYNTHESI
E11	31		TRANSFORMING GROWTH FACTOR ALPHA --BLOOD --BL
E12	3		TRANSFORMING GROWTH FACTOR ALPHA --CEREBROSPIN

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Ref	Items	Index-term
E13	2	TRANSFORMING GROWTH FACTOR ALPHA --CHEMICAL SY
E14	85	TRANSFORMING GROWTH FACTOR ALPHA --CHEMISTRY -
E15	16	TRANSFORMING GROWTH FACTOR ALPHA --DEFICIENCY
E16	1	TRANSFORMING GROWTH FACTOR ALPHA --DIAGNOSTIC
E17	25	TRANSFORMING GROWTH FACTOR ALPHA --DRUG EFFECT
E18	694	TRANSFORMING GROWTH FACTOR ALPHA --GENETICS --
E19	85	TRANSFORMING GROWTH FACTOR ALPHA --IMMUNOLOGY
E20	16	TRANSFORMING GROWTH FACTOR ALPHA --ISOLATION A
E21	600	TRANSFORMING GROWTH FACTOR ALPHA --METABOLISM
E22	4	TRANSFORMING GROWTH FACTOR ALPHA --PHARMACOKIN
E23	760	TRANSFORMING GROWTH FACTOR ALPHA --PHARMACOLOG
E24	389	TRANSFORMING GROWTH FACTOR ALPHA --PHYSIOLOGY

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E25	2		TRANSFORMING GROWTH FACTOR ALPHA --RADIATION E
E26	40		TRANSFORMING GROWTH FACTOR ALPHA --SECRETION -
E27	15		TRANSFORMING GROWTH FACTOR ALPHA --THERAPEUTIC
E28	7		TRANSFORMING GROWTH FACTOR ALPHA --TOXICITY --
E29	10		TRANSFORMING GROWTH FACTOR ALPHA --URINE --UR
E30	0	1	TRANSFORMING GROWTH FACTOR ALPHA RECEPTOR
E31	10		TRANSFORMING GROWTH FACTOR ALPHA-PSEUDOMONAS E
E32	14501	9	TRANSFORMING GROWTH FACTOR BETA
E33	216		TRANSFORMING GROWTH FACTOR BETA --ADMINISTRATI
E34	19		TRANSFORMING GROWTH FACTOR BETA --ADVERSE EFFE
E35	7		TRANSFORMING GROWTH FACTOR BETA --AGONISTS --A
E36	912		TRANSFORMING GROWTH FACTOR BETA --ANALYSIS --A

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E37	435	TRANSFORMING GROWTH FACTOR BETA --ANTAGONISTS
E38	1614	TRANSFORMING GROWTH FACTOR BETA --BIOSYNTHESIS
E39	459	TRANSFORMING GROWTH FACTOR BETA --BLOOD --BL
E40	35	TRANSFORMING GROWTH FACTOR BETA --CEREBROSPINA
E41	1	TRANSFORMING GROWTH FACTOR BETA --CHEMICAL SYN
E42	231	TRANSFORMING GROWTH FACTOR BETA --CHEMISTRY --
E43	41	TRANSFORMING GROWTH FACTOR BETA --CLASSIFICATI
E44	41	TRANSFORMING GROWTH FACTOR BETA --DEFICIENCY -
E45	2	TRANSFORMING GROWTH FACTOR BETA --DIAGNOSTIC U

E46 137 TRANSFORMING GROWTH FACTOR BETA --DRUG EFFECTS
 E47 3100 TRANSFORMING GROWTH FACTOR BETA --GENETICS --G
 E48 1 TRANSFORMING GROWTH FACTOR BETA --HISTORY --HI

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Ref Items Index-term
 E49 693 TRANSFORMING GROWTH FACTOR BETA --IMMUNOLOGY -
 E50 92 TRANSFORMING GROWTH FACTOR BETA --ISOLATION AN

?pp

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?s e32-e50

14501 TRANSFORMING GROWTH FACTOR BETA
 216 TRANSFORMING GROWTH FACTOR BETA --ADMINISTRATI
 19 TRANSFORMING GROWTH FACTOR BETA --ADVERSE EFFE
 7 TRANSFORMING GROWTH FACTOR BETA --AGONISTS --A
 912 TRANSFORMING GROWTH FACTOR BETA --ANALYSIS --A
 435 TRANSFORMING GROWTH FACTOR BETA --ANTAGONISTS
 1614 TRANSFORMING GROWTH FACTOR BETA --BIOSYNTHESIS
 459 TRANSFORMING GROWTH FACTOR BETA --BLOOD --BL
 35 TRANSFORMING GROWTH FACTOR BETA --CEREBROSPINA
 1 TRANSFORMING GROWTH FACTOR BETA --CHEMICAL SYN
 231 TRANSFORMING GROWTH FACTOR BETA --CHEMISTRY --
 41 TRANSFORMING GROWTH FACTOR BETA --CLASSIFICATI
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 137 TRANSFORMING GROWTH FACTOR BETA --DRUG EFFECTS
 3100 TRANSFORMING GROWTH FACTOR BETA --GENETICS --G
 1 TRANSFORMING GROWTH FACTOR BETA --HISTORY --HI
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 92 TRANSFORMING GROWTH FACTOR BETA --ISOLATION AN

S1 14501 E32-E50

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R3	14501	X		DC=D12.644.900.720. (TRANSFORMING GROWTH FACTOR BETA)
R4	14501	X		DC=D24.185.348.900.720. (TRANSFORMING GROWTH FACTOR BETA)
R5	14501	X		DC=D24.611.350.400.800. (TRANSFORMING GROWTH FACTOR BETA)
R6	0	X	1	BONE-DERIVED TRANSFORMING GROWTH FACTOR
R7	0	X	1	PLATELET TRANSFORMING GROWTH FACTOR
R8	1	X	1	TGF-BETA
R9	19034	B	102	GROWTH SUBSTANCES
R10	2125	B	7	TRANSFORMING GROWTH FACTORS

?s r1-r8

14501 TRANSFORMING GROWTH FACTOR BETA
 14501 DC=D11.303.900.720. (TRANSFORMING GROWTH FACTOR BETA)
 14501 DC=D12.644.900.720. (TRANSFORMING GROWTH FACTOR BETA)
 14501 DC=D24.185.348.900.720. (TRANSFORMING GROWTH FACTOR BETA)
 14501 DC=D24.611.350.400.800. (TRANSFORMING GROWTH FACTOR BETA)
 0 BONE-DERIVED TRANSFORMING GROWTH FACTOR
 0 PLATELET TRANSFORMING GROWTH FACTOR
 1 TGF-BETA

S2 14501 R1-R8

?e r8

Ref	Items	Type	RT	Index-term
R1	1		1	*TGF-BETA
R2	14501	X	9	TRANSFORMING GROWTH FACTOR BETA

?s r1-r2

1 TGF-BETA
 14501 TRANSFORMING GROWTH FACTOR BETA

S3 14501 R1-R2

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Set	Items	Description
S1	14501	E32-E50
S2	14501	R1-R8
S3	14501	R1-R2
?s (s1 or s2 or s3) and b4		
	14501	S1
	14501	S2
	14501	S3
	8199	B4

Set	Items	Description
S1	14501	E32-E50
S2	14501	R1-R8
S3	14501	R1-R2
S4	15	(S1 OR S2 OR S3) AND B4
S5	197	(S1 OR S2 OR S3) (3N) (BETA? (2N) 4)
S6	197	(S1 OR S2 OR S3) (3N) (BETA (2N) 4) NOT L4
S7	839	(S1 OR S2 OR S3) (N) (BETA)
S8	0	(S1 OR S2 OR S3) (N) (BETA (N) 4)
S9	0	(S1 OR S2 OR S3) (N) (BETA (2N) 4)
S10	0	(S1 OR S2 OR S3) (N) (B (2N) 4)
S11	2	FACTOR (N) BETA4

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11/9/1

DIALOG(R)File 155:MEDLINE(R)

12782530 21454090 PMID: 11567773

Adjuvant effects of IL-1beta, IL-2, IL-8, IL-15, IFN-alpha, IFN-gamma TGF-beta4 and lymphotactin on DNA vaccination against Eimeria acervulina.

Min W; Lillehoj H S; Burnside J; Weining K C; Staeheli P; Zhu J J
Parasite Biology, Epidemiology, Systematics Laboratory, Animal and Natural Resources Institute, BARC-East, Building 1040, US Department of Agriculture, Beltsville, MD 20705, USA.

Vaccine (England) Oct 12 2001, 20 (1-2) p267-74, ISSN 0264-410X
Journal Code: 8406899

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

Eight chicken cytokine genes (IL-1beta, IL-2, IL-8, IL-15, IFN-alpha, IFN-gamma, TGF-beta4, lymphotactin) were evaluated for their adjuvant effect on a suboptimal dose of an Eimeria DNA vaccine carrying the 3-1E parasite gene (pcDNA3-1E). Chickens were given two subcutaneous injections with 50 microg of the pcDNA3-1E vaccine plus a cytokine expression plasmid 2 weeks apart and challenged with Eimeria acervulina 1 week later. IFN-alpha (1 microg) or 10 microg of lymphotactin expressing plasmids, when given simultaneously with the pcDNA3-1E vaccine, significantly protected against body weight loss induced by E. acervulina. Parasite replication was significantly reduced in chickens given the pcDNA3-1E vaccine along with 10 microg of the IL-8, lymphotactin, IFN-gamma, IL-15, TGF-beta4, or IL-1beta plasmids compared with chickens given the pcDNA3-1E vaccine alone. Flow cytometric analysis of duodenum intraepithelial lymphocytes showed chickens that received the pcDNA3-1E vaccine simultaneously with the IL-8 or IL-15 genes had significantly increased CD3+ cells compared with vaccination using pcDNA3-1E alone or in combination with the other cytokine genes tested. These results indicate that the type and the dose of cytokine genes injected into chickens influence the quality of the local immune response to DNA vaccination against coccidiosis.

Tags: Animal; Comparative Study; Support, U.S. Gov't, Non-P.H.S.

Descriptors: *Adjuvants, Immunologic; *Coccidiosis--veterinary--VE; *Eimeria--immunology--IM; *Interferons--immunology--IM; *Interleukins--immunology--IM; *Lymphokines--immunology--IM; *Poultry Diseases--prevention and control--PC; *Sialoglycoproteins--immunology--IM; *Transforming Growth Factor beta--immunology--IM; Chickens; Coccidiosis--immunology--IM; Coccidiosis--prevention and control--PC; Drug Evaluation, Preclinical; Duodenum--immunology--IM; Duodenum--parasitology--PS; Genetic Vectors--administration and dosage--AD; Genetic Vectors--genetics--GE; Interferon Type II--genetics--GE; Interferon Type II--immunology--IM; Interferon-alpha--genetics--GE; Interferon-alpha--immunology--IM; Interferons--genetics--GE; Interleukin-1--genetics--GE; Interleukin-1--immunology--IM; Interleukin-15--genetics--GE; Interleukin-15--immunology--IM; Interleukin-2--genetics--GE; Interleukin-2--immunology--IM; Interleukin-8--genetics--GE; Interleukin-8--immunology--IM; Interleukins--genetics--GE; Lymphokines--genetics--GE; Parasite Egg Count; Poultry Diseases--immunology--IM; Sialoglycoproteins--genetics--GE; Specific Pathogen-Free Organisms; Transforming Growth Factor beta--genetics--GE; Vaccination--veterinary--VE; Vaccines, DNA--genetics--GE; Vaccines,

DNA--immunology--IM; Weight Gain

CAS Registry No.: 0 (Adjuvants, Immunologic); 0 (Genetic Vectors); 0 (Interferon-alpha); 0 (Interleukin-1); 0 (Interleukin-15); 0 (Interleukin-2); 0 (Interleukin-8); 0 (Interleukins); 0 (Lymphokines); 0 (Sialoglycoproteins); 0 (Transforming Growth Factor beta); 0 (Vaccines, DNA); 0 (lymphotactin); 0 (transforming growth factor beta4); 82115-62-6 (Interferon Type II); 9008-11-1 (Interferons)
Record Date Created: 20010924

11/9/2

DIALOG(R) File 155:MEDLINE(R)

09908583 98328280 PMID: 9665343

Temporal and site-specific expression of transforming growth factor - beta4 in human endometrium.

Tabibzadeh S; Lessey B; Satyaswaroop P G

Department of Pathology, Moffitt Cancer Center, Tampa, FL 33612, USA.
tabibzadeh@bioscience.org

Molecular human reproduction (ENGLAND) Jun 1998, 4 (6) p595-602,
ISSN 1360-9947 Journal Code: 9513710

Contract/Grant No.: CA46866; CA; NCI; CA62211; CA; NCI; HD34824; HD; NICHD

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

We recently identified a novel member of the transforming growth factor (TGF)-beta superfamily and showed that this gene, designated as endometrial bleeding associated factor (ebaf), or TGFbeta4, has a unique expression pattern in human endometrium. By Northern blot analysis, we showed that this gene was expressed in human endometrium during the late secretory and menstrual phases and was absent in proliferative, early and mid-secretory endometria. In this report, we show by in-situ hybridization that the mRNA of the TGF-beta4 is not expressed in the proliferative endometria. On the other hand, focal expression of the TGFbeta4 mRNA first appears in some endometrial glands in the mid-secretory phase. The TGFbeta4 mRNA is strongly expressed in the endometrial stroma during the late secretory and menstrual phases of the cycle. We raised a polyclonal rabbit antiserum against a peptide at the C terminal of the protein. Western blot analysis using affinity purified antiserum shows that the TGFbeta4 precursor detected in the endometrium as well as placenta is 41 kDa. Bands in the range of 45-51 kDa are also present in human endometrium, more predominantly during the late secretory phase. Immunohistochemical staining shows a low level of immunoreactivity for TGFbeta4 in the early, mid- and late proliferative and early and mid-secretory endometria. A strong immunoreactivity for TGFbeta4 is present in the stroma and to lesser extent in the endometrial glands in late secretory and menstrual endometria. The specificity of staining was shown by neutralizing the activity of the antibody with the synthetic peptide used for raising the antibody and by omitting the antibody. The findings show that TGFbeta4, both at the mRNA and protein levels, exhibits temporal and site specific expression in human endometrium.

Tags: Animal; Female; Human; Support, Non-U.S. Gov't; Support, U.S. Gov't, P.H.S.

Descriptors: *Endometrium--metabolism--ME; *Gene Expression Regulation; *Menstrual Cycle--genetics--GE; *Transforming Growth Factor beta --biosynthesis--BI; Amino Acid Sequence; Blotting, Western; Immunoenzyme Techniques; In Situ Hybridization; Molecular Sequence Data; Multigene Family; Peptide Fragments--immunology--IM; RNA, Messenger--biosynthesis--BI; RNA, Messenger--genetics--GE; Rabbits; Transforming Growth Factor beta --genetics--GE; Transforming Growth Factor beta--immunology--IM

CAS Registry No.: 0 (Peptide Fragments); 0 (RNA, Messenger); 0 (Transforming Growth Factor beta)

Record Date Created: 19981022

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\$0.42 2 Type(s) in Format 9
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